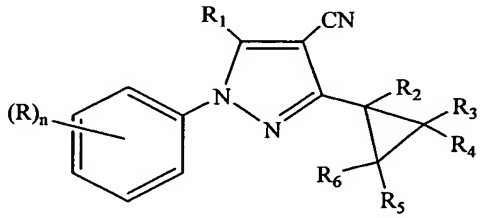
What is claim d is:

1. A composition for prevention, amelioration or control of external parasites on animals and humans comprising a pharmaceutically acceptable carrier and an ectoparasiticidally effective amount of a compound of formula I



(I)

or a pharmaceutically acceptable salt thereof wherein

R is halogen, OR_7, SO_mR_8, NO_2, CN, C_1-C_6haloalkyl or an optionally substituted C_1-C_6alkyl group;

n is 0 or an integer of 1, 2 or 3;

m is 0 or an integer of 1 or 2;

R_1 is H, halogen, NO_2, NR_9R_{10}, $NR_{11}COR_{12}$, $NCHNR_9R_{10}$ or $NCHOR_{13}$;

R_2, R_3, R_4, R_5 and R_6 are each independently H, halogen or a C_1-C_4alkyl, aryl or heteroaryl group each optionally substituted;

R_7 is H or a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group each optionally substituted;

R_8 is a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group, each optionally substituted;

R_9 and R_{10} are each independently H, C_1-C_4haloalkyl or a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group each optionally substituted or R_9 and R_{10} may be taken together with the atom to which they are attached to form a 5- to 7-membered ring optionally containing 1 or 2 additional heteroatoms selected from O, N or S;

R_{11} is H, COR_{12} or an optionally substituted C_1-C_4alkyl group;

R_{12} is a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group each optionally substituted; and

R_{13} is H or a C_1-C_6alkyl, aryl or heteroaryl group each optionally substituted;

or a stereoisomer or tautomer thereof.

2. The composition according to claim 1 wherein formula I has the proviso that R_3, R_4, R_5 and R_6 are not all –H, unless R_1 is halogen.

3. The composition according to claim 2 wherein R is halogen or haloalkyl.

4. The composition according to claim 2 wherein R_1 is H, halogen or NR_9R_{10}.

5. The composition according to claim 1 wherein R_5 and R_6 are H.

6. The composition according to claim 3 wherein R_2 is H, halogen, methyl or an optionally substituted phenyl group.

7. The composition according to claim 6 wherein R_1 is H or Cl.

8. The composition according to claim 7 wherein R is halogen or CF_3 and n is 3.

9. The composition according to claim 8 wherein R_2 is Cl or methyl and R_3 and R_4 are each independently H, Cl or Br.

10. The composition according to claim 2 wherein said compound is selected from the group consisting of:

5-chloro-3-(2,2-dichloro-1methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1,3-dimethylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

3-(2,2-dibromo-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

5-chloro-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;

5-amino-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;

5-bromo-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

5-amino-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;

5-chloro-3-cyclopropyl-1--[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;

5-chloro-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;

5-bromo-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;

5-bromo-3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-nitro-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-iodo-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-(dimethylamino)-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopyrazol)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-(diethylamino)-1H-pyrazole-4-carbonitrile;

5-[(cyclopropanecarbonyl)amino-]-3-(2,2-dichloro-1methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

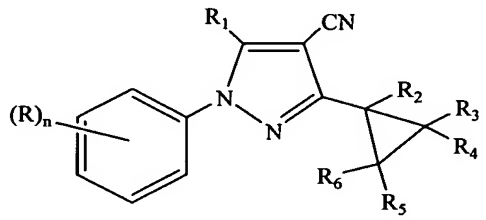
N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2H-pyrazol-3-yl}-formimidic acid methyl ester;

N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2-H-pyrazol-3-yl}-formimidic acid propyl ester;

N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2H-pyrazol-3-yl}-formimidic acid ethyl ester;

the stereoisomers thereof; and the tautomers thereof; or a pharmaceutically acceptable salt thereof.

11. A method for the prevention, amelioration or control of ectoparasitic infection or infestation in a homeothermic animal which comprises providing to a homeothermic animal in need thereof a prophylactically, therapeutically or pharmaceutically effective amount of a compound of formula I



(I)

or a pharmaceutically acceptable salt thereof wherein

R is halogen, OR_7, SO_mR_8, NO_2, CN, C_1-C_6haloalkyl or an optionally substituted C_1-C_6alkyl group;

n is 0 or an integer of 1, 2 or 3;

m is 0 or an integer of 1 or 2;

R_1 is H, halogen, NO_2, NR_9R_{10}, $NR_{11}COR_{12}$, $NCHNR_9R_{10}$ or $NCHOR_{13}$;

R_2, R_3, R_4, R_5 and R_6 are each independently H, halogen or a C_1-C_4alkyl, aryl or heteroaryl group each optionally substituted;

R_7 is H or a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group each optionally substituted;

R_8 is a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group, each optionally substituted;

R_9 and R_{10} are each independently H, C_1-C_4haloalkyl or a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group each optionally substituted or R_9 and R_{10} may be taken together with the atom to

which they are attached to form a 5- to 7-membered ring optionally containing 1 or 2 additional heteroatoms selected from O, N or S;

R_{11} is H, COR_{12} or an optionally substituted C_1-C_4alkyl group;

R_{12} is a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group each optionally substituted; and

R_{13} is H or a C_1-C_6alkyl, aryl or heteroaryl group each optionally substituted; or a stereoisomer or tautomer thereof.

12. The method according to claim 11 wherein the formula I has the proviso that R_3, R_4, R_5 and R_6 are not all –H, unless R_1 is halogen.

13. The method according to claim 12 wherein R is halogen or haloalkyl and n is 3.

14. The method according to claim 13 wherein R_5 and R_6 are H.

15. The method according to claim 14 wherein R_2 is H, halogen, methyl or an optionally substituted phenyl group.

16. The method according to claim 12 wherein the ectoparasite is selected from the group consisting of Diptera; Muscidae; Acarina; and Siphonáptera.

17. The method according to claim 16 wherein the ectoparasite is selected from the group consisting of fleas; ticks; lice; blow flies; face flies and horn flies.

18. The method according to claim 16 wherein the homeothermic animal is selected from the group consisting of cattle; sheep; horse; goat; pig; camel; water buffalo; donkey; rabbit; fallow deer; reindeer; mink; chinchilla; raccoon; chicken; geese; turkey; duck; dog and cat.

19. The method according to claim 17 wherein the homeothermic animal is selected from the group consisting of cattle, sheep, horse, dog, and cat.

20. A veterinary pour-on composition which comprises: a spreading oil; an aliphatic or aromatic hydrocarbon, mono or polyhydric alcohol, a C_1-C_{10} alkyl

ketone, or a mixture thereof; and an ectoparasiticidally effective amount of a compound of formula I according to claim 1.

21. The composition according to claim 20 wherein formula I has the proviso that R_3, R_4, R_5 and R_6 are not all –H, unless R_1 is halogen.

22. The composition according to claim 21 wherein R is halogen or haloalkyl and n is 3.

23. The composition according to claim 22 wherein said compound is selected from the group consisting of:

5-chloro-3-(2,2-dichloro-1methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1,3-dimethylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

3-(2,2-dibromo-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

5-chloro-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;

5-amino-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;

5-bromo-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

5-amino-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;

5-chloro-3-cyclopropyl-1--[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;
5-chloro-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
5-bromo-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
5-bromo-3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-nitro-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-iodo-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-(dimethylamino)-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopyrazol)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-(diethylamino)-1H-pyrazole-4-carbonitrile;
5-[(cyclopropanecarbonyl)amino-]-3-(2,2-dichloro-1methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2H-pyrazol-3-yl}-formimidic acid methyl ester;
N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2-H-pyrazol-3-yl}-formimidic acid propyl ester;
N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2-H-pyrazol-3-yl}-formimidic acid ethyl ester;
the stereoisomers thereof; and the tautomers thereof.

24. A veterinary pour-on composition which comprises: approximately 40-50% by weight xylene; approximately 20-30% by weight cyclohexanone; approximately 5-15% vegetable or mineral oil or a combination thereof; and approximately 10-25% of a compound selected from the group consisting of:
5-chloro-3-(2,2-dichloro-1methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1,3-dimethylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
3-(2,2-dibromo-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
5-chloro-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;
5-amino-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;
5-bromo-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
5-amino-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
5-chloro-3-cyclopropyl-1--[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;
5-chloro-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
5-bromo-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
5-bromo-3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-nitro-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-iodo-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-(dimethylamino)-1H-pyrazole-4-carbonitrile;

3-(2,2-dichloro-1-methylcyclopyrazol)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-(diethylamino)-1H-pyrazole-4-carbonitrile;
5-[(cyclopropanecarbonyl)amino-]-3-(2,2-dichloro-1methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2H-pyrazol-3-yl}-formimidic acid methyl ester;
N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2-H-pyrazol-3-yl}-formimidic acid propyl ester;
N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2-H-pyrazol-3-yl}-formimidic acid ethyl ester;
the stereoisomers thereof; and the tautomers thereof.

25. The composition according to claim 24 wherein an effective dosage of said compound is within the range of about 0.1 mg/kg to 100 mg/kg of animal body weight.

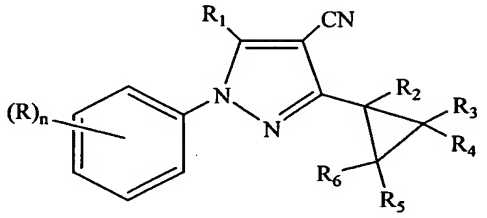
26. A veterinary composition which comprises a pharmaceutically acceptable carrier and about 0.1 ppm to 5000 ppm of a compound selected from the group consisting of:
5-chloro-3-(2,2-dichloro-1methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1,3-dimethylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
3-(2,2-dibromo-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;

5-chloro-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;
5-amino-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;
5-bromo-3-(2,2-dibromo-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
5-amino-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
5-chloro-3-cyclopropyl-1--[2,6-dichloro-4-(trifluoromethyl)phenyl-1H-pyrazole-4-carbonitrile;
5-chloro-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
5-bromo-3-(2,2-dichloro-1-methylcyclopropyl)-1-(2,4,6-trichlorophenyl)-1H-pyrazole-4-carbonitrile;
5-bromo-3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-nitro-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-iodo-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-(dimethylamino)-1H-pyrazole-4-carbonitrile;
3-(2,2-dichloro-1-methylcyclopyrazol)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-5-(diethylamino)-1H-pyrazole-4-carbonitrile;
5-[(cyclopropanecarbonyl)amino-]-3-(2,2-dichloro-1methylcyclopropyl)-1-[2,6-dichloro-4-(trifluoromethyl)phenyl]-1H-pyrazole-4-carbonitrile;
N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2H-pyrazol-3-yl}-formimidic acid methyl ester;
N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2-H-pyrazol-3-yl}-formimidic acid propyl ester;
N-{4-cyano-5-(2,2-dichloro-1-methyl-cyclopropyl)-2-[2,6-dichloro-4-(trifluoromethyl)-phenyl]-2-H-pyrazol-3-yl}-formimidic acid ethyl ester;
the stereoisomers thereof; and the tautomers thereof.

27. The composition according to claim 26 which comprises about 0.5 ppm to 1000 ppm of said compound.

28. The composition according to claim 27 which comprises about 0.2 ppm to 20 ppm of said compound.

29. A compound of formula I



(I)

or a pharmaceutically acceptable salt thereof wherein

R is halogen, OR_7, SO_mR_8, NO_2, CN, C_1-C_6haloalkyl or an optionally substituted C_1-C_6alkyl group;

n is 0 or an integer of 1, 2 or 3;

m is 0 or an integer of 1 or 2;

R_1 is H, halogen, NO_2, NR_9R_{10}, $NR_{11}COR_{12}$, $NCHNR_9R_{10}$ or $NCHOR_{13}$;

R_2, R_3, R_4, R_5 and R_6 are each independently H, halogen or a C_1-C_4alkyl, aryl or heteroaryl group each optionally substituted;

R_7 is H or a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group each optionally substituted;

R_8 is a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group, each optionally substituted;

R_9 and R_{10} are each independently H, C_1-C_4haloalkyl or a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group each optionally substituted or R_9 and R_{10} may be taken together with the atom to which they are attached to form a 5- to 7-membered ring optionally containing 1 or 2 additional heteroatoms selected from O, N or S;

R_{11} is H, COR_{12} or an optionally substituted C_1-C_4alkyl group;

R_{12} is a C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, aryl or heteroaryl group each optionally substituted; and

R_{13} is H or a C_1-C_6alkyl, aryl or heteroaryl group each optionally substituted; or a stereoisomer or tautomer thereof.

30. The compound of claim 29 wherein formula I has the proviso that R_3, R_4, R_5 and R_6 are not all –H, unless R_1 is halogen.